

PATENT APPLICATION

**Species Specific Identification Of Spore-Producing Microbes Using The
Gene Sequence Of Small Acid Soluble Spore Coat Proteins For
Amplification Based Diagnostics**

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CROSS-REFERENCES TO RELATED APPLICATIONS

[01] The present application is a Continuation-In-Part application ("CIP") of U.S. Application Serial No. 09/590,759, filed June 8, 2000, which claims priority benefit of U.S. Provisional Application No. 60/138,167, filed on June 8, 1999, and U.S. Provisional Application No. 60/192,206, filed on March 27, 2000. The aforementioned applications are explicitly incorporated herein by reference in their entirety and for all purposes.

STATEMENT AS TO RIGHTS TO INVENTIONS MADE UNDER
FEDERALLY SPONSORED RESEARCH OR DEVELOPMENT

[02] This invention was made under work supported by the U.S. Department of Energy under DOE Contract No.: DE-AC03-76SF00098. The government has certain rights in this invention.

REFERENCE TO A "SEQUENCE LISTING," A TABLE, OR A COMPUTER
PROGRAM LISTING APPENDIX SUBMITTED ON A COMPACT DISK.

[03] NOT APPLICABLE

BACKGROUND OF THE INVENTION

Field of the Invention

[04] The present invention relates to methods and compositions for the detection of *Bacillus* species such as *Bacillus anthracis* and *Bacillus globigii* as well as *Clostridium perfringens*. It relies on nucleic acid sequence differences in spore protein genes carried in the genomic sequence of these organisms.

[05] The genus *Bacillus* is composed of rod-shaped, gram-positive, aerobic or (under some conditions) anaerobic bacteria widely found in soil and water. Most strains of *Bacillus* are not pathogenic for humans and only infect them incidentally in their role as soil organisms; a notable exception is *Bacillus anthracis*, which causes anthrax in humans and domestic animals. In addition to its role as a naturally occurring pathogen, *Bacillus anthracis*

may also be used as a biological weapon. Because *Bacillus* organisms are widely distributed in the environment, and because they are very closely related genetically, there is need for a reliable method to distinguish species members in various types of samples.

[06] Considerable efforts have been made to develop sensitive, species specific tests for these *Bacillus* microorganisms with only limited results; examples of such efforts follow:

[07] Makino *et al.* describes a PCR-based test using primers to the cap region of plasmid pXO2 (codes for a protein essential for capsulation) which was amplified by PCR and probed with the *B. anthracis* capA DNA, Makino *et al.*, "Direct Detection of *Bacillus anthracis* DNA in animals by Polymerase Chain Reaction", *J. Clin. Micro.* 1993 March 31(3):547-51.

[08] Keim *et al.* describes an AFLP (amplified-fragment length polymorphism) technique employing high resolution electrophoresis to examine restriction fragments of PCR products, Keim, P; Kalif, A; Schupp, J; Hill, K; Travis, SE; Richmond, K; Adair, DM; Hugh-Jones, M; Kuske, CR; Jackson, P., "Molecular evolution and diversity in *Bacillus anthracis* as detected by amplified fragment length polymorphism markers, *Journal of Bacteriology*, 1997 Feb, 179(3):818-24.

[09] Similarly, Andersen *et al.* describes a technique using arbitrarily primed (AP)-PCR "fingerprints" that detect variable numbers of repeats in different samples, Andersen *et al.* "Identification of a region of genetic variability among *Bacillus anthracis* strains and related species," *J. Bacteriology*, 1996 Jan, 178(2):377-84; Beyer, W; Glöckner, P; Otto, J; Böhm, R. "A nested PCR method for the detection of *Bacillus anthracis* in environmental samples collected from former tannery sites" *Microbiological Research*, 1995 May, 150(2):179-86.

[10] Reif *et al.* describes a PCR technique for identifying spores of *B. anthracis*. Primers specific for the capB region of plasmid pXO2 are used, Reif *et al.* "Identification of capsule forming *Bacillus anthracis* spores with the PCR and a novel dual- probe hybridization format," *Applied and Environmental Microbiology*, 1994 May, 60(5):1622-5.

[11] Brightwell *et al.* describes a PCR reaction for detecting the presence of *B. anthracis* plasmid pXO2, Brightwell *et al.* "Development of internal controls for PCR detection of *Bacillus anthracis*" *Molecular and Cellular Probes*, (1998) 12, 367-377.

[12] Jackson *et al.* describes PCR analysis using primers that detect the *vrrA* gene variable region on the *B. anthracis* chromosome, Jackson *et al.*, *P.N.A.S.* 95:1224-1229,

February 1998, "PCR analysis of tissue samples from the 1979 Sverdlovsk anthrax victims: the presence of multiple *Bacillus anthracis* strains in different victims."

[13] Patral *et al.* describes a 277-bp long noncoding DNA fragment, Ba813, that was isolated from an avirulent *Bacillus anthracis* strain 7700 genomic library. Two oligonucleotides derived from the Ba813 sequence were used as primers in polymerase chain reaction tests on genomic DNA from 28 *Bacillus anthracis* and from 33 heterologous bacteria strains. A specific, 152-bp long DNA fragment was amplified only when *Bacillus anthracis* DNA was used as the target [Note: primers eventually demonstrated non-specific (McKinney, manuscript in preparation)], Patra, G; Sylvestre, P; Ramisse, V; Thaerasse, J; Guesdon, JL."Isolation of a specific chromosomal DNA sequence of *Bacillus anthracis* and its possible use in diagnosis," *Fems Immunology and Medical Microbiology*, 1996 Oct, 15 4 :223-31.

[14] Most of the *Bacillus anthracis* PCR detection systems have targeted plasmid sequences, yet mobility of these genetic elements make them an unreliable detection target. (Battisti, L; Green, BD; Thorne, CB. Mating system for transfer of plasmids among *Bacillus anthracis*, *Bacillus cereus*, and *Bacillus thuringiensis*. *Journal of Bacteriology*, 1985 May, 162(2):543-50. Green, BD; Battisti, L; Thorne, CB. Involvement of Tn4430 in transfer of *Bacillus anthracis* plasmids mediated by *Bacillus thuringiensis* plasmid pXO12. *Journal of Bacteriology*, 1989 Jan, 171(1):104-13) Several of the PCR type assays to genomic sequences target non-gene-coding sequences, which often proves not to correlate well with species identity. The weaknesses in each of these systems underscores the need for a more reliable *Bacillus* identification method.

SUMMARY OF THE INVENTION

[15] It is an object of the present invention to provide methods and materials for specifically detecting and identifying members of the *Bacillus* family, especially *Bacillus anthracis*, to the exclusion of other members of the genus *Bacillus*, which are closely related. These methods and materials rely on the discovery that certain small acid-soluble protein (sasp) gene targets are present in the bacterial coats of *Bacillus* and *Clostridium* organisms. These sasp gene targets may be identified by a technique described here as "heterologous PCR". More specifically, these methods and materials provide means for specifically identifying certain members of the *Bacillus* genus, especially *Bacillus anthracis* and *Bacillus globigii*, and members of the *Clostridium* genus, especially *Clostridium perfringens*. Using heterologous PCR, novel gene targets are identified that yield species-specific regions for detection and identification. Based on this technique, assay techniques for identifying the

organisms *Bacillus anthracis*, *Bacillus globigii* and *Clostridium perfringens* have been developed.

[16] Another aspect of the invention comprises use of a small acid soluble protein (sasp-B) gene sequence in *Bacillus anthracis* that contains a specific region of stable nucleotide insertion absent in even the closest *Bacillus* relatives. This unique biomarker is a probe-binding region for the specific detection of *Bacillus anthracis*. Additional regions of sequence within the gene were used to develop a *Bacillus anthracis* specific amplification and detection system. The system is capable of distinguishing *Bacillus anthracis* from near neighbors during two phases of the amplified assay: during the amplification reaction, and by probe hybridization to the *Bacillus anthracis* specific biomarker within the amplified product for sequence confirmation.

[17] As another aspect of the invention, the present *Bacillus anthracis* detection system either alone, or multiplexed with one or more additional genomic and/or virulence plasmid markers, is of use as a diagnostic or confirmatory tool in Public Health and clinical laboratories, as well as in the law enforcement sector. The system may be adapted to a variety of amplification and detection platforms in order to accommodate the technical and fiscal capabilities of the laboratory, or used in 'field' settings.

[18] As yet another aspect of the invention, building upon the discovery that small acid-soluble spore protein genes maintain species-specific sequence signatures, analogous regions were identified among the sasp of *Bacillus globigii* (which is used as a non-pathogenic 'surrogate' for *Bacillus anthracis* in research and development applications, defense, and emergency response modeling, etc.). Taking advantage of this discovery, *Bacillus globigii* specific PCR was designed and reduced to practice.

[19] Finally, using the same principals and strategy, primers were designed to a sasp gene sequence of *Clostridium perfringens*, (an acknowledged biological weapon) which was successfully amplified by the system.

[20] Another aspect of the invention is to provide a two-step amplification/detection system. A particular sasp gene is selected for amplification, and its identity determined by a species-specific probe. Although the present invention is described in detail in connection with a PCR, it is understood that, based on the present teachings, other amplification and/or identification systems could be devised.

BRIEF DESCRIPTION OF THE DRAWINGS

[21] Figure 1. is a multiple ClustalW DNA Sequence Alignment of sasp-B Amplicons from 38 *Bacillus anthracis* strains (Seq. ID No. 13 through 50). Bases 1 - 90 are in Fig. 1A, 91 - 180 are in Fig. 1-B, and 181 - 240 are in 1C.

[22] Figure 2. is a ClustalW multiple sasp-B DNA Sequence Alignment of *Bacillus anthracis*, *Bacillus thuringiensis* and *Bacillus cereus* strains (Seq. ID No. 51 through 87). Bases 1 - 90 are in Fig. 2A, 91 - 180 are in Fig. 2-B, and 181 - 240 are in 2C.

[23] Figure 3. is a representation of *Bacillus globigii* specific PCR primers targeting Bg sasp-gamma.

DETAILED DESCRIPTION OF THE INVENTION

[24] Described herein are regions of genomic sequence with patterns unique to the target organisms (*Bacillus anthracis*, *Bacillus globigii* and *Clostridium perfringens*) from which primers and probes were designed for specific amplification of target organism DNA, and where feasible, confirmation of amplicon sequence by probe hybridization. Spore coat and spore structural genes were studied because their products are intimately linked with the organism's environmental niche and, phenotype, and therefore distinct identity of each species. In the case of *Bacillus anthracis*, very little genomic sequence data is available, hence published sequence listings from closely related species were used as a starting point from which primers for amplification of *Bacillus anthracis* DNA (a process commonly known as 'heterologous PCR') were designed. Heterologous PCR, then, means PCR using primers known to hybridize to one target to amplify, under conditions of low stringency, another target, in this case, from another species of unknown sequence in the target gene. Some 27 spore genes were screened via heterologous PCR and scores of reaction products sequenced before a sufficiently definitive region of sequence was identified for anthracis specific primer and probe design. The signature which satisfied the specificity criteria, and which is the key to this invention, was found within the coding sequence of the sasp spore structural protein.

Example 1: Database Search & Primer Design Example (*Bacillus anthracis* primer/probe design)

[25] Public databases GenBank and European Molecular Biology Laboratory (EMBL) were queried for "small acid-soluble spore protein" (sasp) DNA sequences. Three *Bacillus cereus* small acid-soluble protein genes were selected from GenBank for

consideration (*Bacillus cereus* being one of the closest relatives to *Bacillus anthracis*).
GenBank accession numbers for these sasp-B DNA sequences are: M13059 for *Bacillus cereus* sasp-1, M13060 for *Bacillus cereus* sasp-2, and M16813 for *Bacillus cereus* sasp-B.

[26] Primer sequences were located within each sasp sequence which would
5 maximize the likelihood of amplifying non-homologous sequences. For instance, whenever
possible the 3' end of a primer was concluded with one or more thymidine residues.
Potential primer sequences were analyzed using Oligo 4.0 primer design software (National
Biosciences, Plymouth, MN) for potential hairpin or concatomers, which might interfere with
hybridization to target DNA. Also using Oligo 4.0 primer design software (National
10 Biosciences, Plymouth, MN), primer sequences were adjusted to match their melting
temperatures as closely as possible to one another, which generally enhances reaction
specificity. The sequence similarity search tool BLAST was queried with the primer
sequences in order to insure that the primers did not recognize any bacterial (or other
microbial) sequences except the targeted *Bacillus* species. Primers were synthesized
15 (Sequence IDs No.1 through 6) using the PerSeptive Biosystems Expedite nucleic acid
synthesis system (Perkin Elmer, Norwalk, Conn.). Oligos were released from columns by
incubation in 29.3% ammonium hydroxide at 55°C. overnight, followed by evaporation of
ammonium hydroxide using the SpeedVac 1SS110 (Savant Corp.). Primers were
resuspended in 10 millimolar tris buffer, pH 8.3, and their concentration measured with a
20 spectrophotometer.

[27] Shown below are primers designed from *Bacillus cereus* sequences for
heterologous PCR and sequencing of *Bacillus anthracis*, as described in Example 2.

B. cereus primers designed for heterologous PCR and sequencing of *B. anthracis*:

Primer name	Sequence (5' to 3')
Bcsasp-B 5'	ATGAGTAAAAACAACAAGGTTAT (SEQ ID NO: 1)
Bcsasp-B 3'	CTGATTTGAGCTAGAAGATTGTGA (SEQ ID NO: 2)
Bcsasp-1 5'	ATGGGAAAAAATAATAGTGGAAGT (SEQ ID NO: 3)
Bcsasp-1 3'	GCGGTTAGCTCTACCAAGT (SEQ ID NO: 4)
Bcsasp-2 5'	ATGTCAGCTAGCACAAATAAATT (SEQ ID NO: 5)
Bcsasp-2 3'	TTATTTTTGGTAACCGCCTAA (SEQ ID NO: 6)

[28] The primers are named according to their corresponding *Bacillus cereus* sasp.

Example 2: Amplification of *Bacillus* species using sasp primers, and analysis of reaction products

5 [29] DNA was prepared according to the method described by Zhou *et al.* (Zhou, J., Bruns, M., and Jiedje, J. 1996. "DNA recovery from soils of diverse composition." *Appl. Environ. Microbiol.* 62:316-322, 1996) from a non-infectious vegetative cells of *Bacillus anthracis* and *Bacillus cereus* type 168 (*Bacillus* Genetic Stock Center). The concentration of the purified DNA was determined by measuring the optical density at 260 nanometer
10 wavelength using the Du 640 spectrophotometer (Beckman, Palo Alto), and by visual comparison to DNA standards of known quantity via agarose gel electrophoresis (Sambrook *et al.* Molecular Cloning, 1989, Cold Spring Harbor Laboratory Press).

[30] For initial trial of the new primers, Tris buffer at pH 8.3, 8.8, and 9.2 was evaluated along with potassium chloride at (final) concentrations of 25 mM and 75 mM and
15 magnesium chloride at (final) concentrations of 1.5 and 3.5 mM were screened using OptiPrime 10X buffers (Stratagene, La Jolla, CA), in order to determine favorable reaction conditions for the new primers. Reaction volumes were 100 microliters and contained approximately 100 nanograms of target DNA (and in the case of negative controls, no DNA).

[31] A GeneAmp 9600 PCR System (Perkin Elmer, Norwalk, Conn) was
20 programmed as follows for thermalcycling of the reaction: A 5 minute 94 degree C. initial denaturation step was followed by 40 three step cycles of 94 degrees C. for 30 sec., 50 degrees C. for 30 sec., 72 degrees C. for 30 sec. A final extension step of 7 minutes at 72 degrees C. completed the thermalcycling.

[32] Amplification products were viewed by ethidium bromide stained
25 nusieve/agarose slab gel electrophoresis. Ten microliters of PCR product was mixed with 2 microliters of 6 X gel loading dye (a filter sterilized solution of 0.25% bromophenol blue plus 40% (wt/vol) sucrose in double distilled sterile water). A molecular weight marker ladder of 100 base pair DNA fragments (Life Technologies, Gaithersburg, MD) was run alongside the amplicons in order to gauge the size of the PCR products. The expected size of reaction
30 products based on the primer locations within the *B. cereus* DNA sequence are: 213 base pairs for sasp-1, 198 base pairs for sasp-2, and 279 base pairs for sasp-B.

[33] Results of this initial experiment are as follows: The major products (as judged by band intensity when viewed by gel) from amplification of *Bacillus anthracis* DNA were the same size as the major products of *Bacillus cereus* amplification using each of the three

primers, regardless of the reaction mixture employed. Thus, it appeared that a *Bacillus anthracis* sasp gene sequence had been obtained.

Example 3: DNA sequence analysis of *Bacillus cereus* sasp-B primed PCR products from

Bacillus anthracis

[34] *Bacillus anthracis* PCR product from Example 2 was separated from primers and other reactants using the QIAquick Gel Extraction kit (Qiagen, Valencia, CA), and sequenced using the Dye Terminator Cycle Sequencing Kit (PE Biosystems, Foster City, CA) using the same *Bacillus cereus* sasp-B primers used to generate the product being sequenced.

Sequencing reactions were performed in a TC-9600 thermalcycler (Perkin Elmer, Norwalk, Conn). The completed sequencing reaction was electrophoresed and the sequence recorded by the Applied Biosystems Prism 377 Automated Sequencer (PE Biosystems, Foster City, CA). Since the PCR products were less than 300 bases, the entire length of each product was sequenced in a single run. The small size also made it possible to check the sequence by alignment of the two strands to one another. After inverting one strand with Gene Jockey sequence conversion software (BioSoft, Cambridge, UK), the strands were aligned using the Baylor College of Medicine Clustal W online multiple sequence alignment utility.

CLUSTAL W is described in Thompson, J.D. Higgins, D.G. Gibson, T.J. (1994) as improving the sensitivity of progressive multiple sequence alignment through sequence weighting, position specific gap penalties and weight matrix choice. *Nucl. Acids Res.* 22:4673-4680. Where there was discrepancy between the two strands (i.e. where the bases did not match in the alignment), electrophoretograms of the sequencing gel run were examined (using the PE Biosystems Editview utility) in order to resolve the discrepancy.

[35] Finished *Bacillus anthracis* sequence was aligned with *Bacillus cereus* published sequence and the differences analyzed. Of greatest interest was the six base region in the *Bacillus anthracis* sasp-B amplicon which was not present in the *Bacillus cereus* published sequence. Alignments of the *Bacillus anthracis* sequence with the published *Bacillus cereus* sequence for the region between each primer pair follow.

[36] In these alignments, dots signify a match with the sequence shown; only mismatches are spelled out, in order to emphasize them. Primer sequences are not included, but would be extensions of the 5' and 3' ends of the sequences shown.

The *Bacillus anthracis* and *B. cereus* sasp-1 sequence alignment did not show significant differences

B.cer 1 CGTAATGAAGTATTAGTTCGAGGCGCTGAACAAGCTCTTGATCAAATGAAATATGAAATT
 B.anth 1T.....T.....
 B.cer 61 GCACAAGAGTTTGGTGTACAACCTGGTGCAGATACAACAGCTCGTTCAAACGGATCTGTT
 B.anth 61T.....
 B.cer 121 GGTGGTGAAATTACAAAACGTTTAGTAGCAATGGCAGAACA (SEQ ID NO: 7)
 B.anth 121T..... (SEQ ID NO: 8)

The *Bacillus anthracis* and *B. cereus* sasp-2 sequence alignment did not show significant differences

B.cer 1 AGCGGTTCTGGTGCTGAATCAGCATTAGACCAAATGAAATACGAAATCGCTCAAGAGTT
 B.anth 1
 B.cer 61 TGGTGTTCAACTGGAGCTGATGCAACAGCTCGCGCTAACGGTTCTGTTGGTGGCGAAAT
 B.anth 61
 B.cer 121 CACTAAACGTCTAGTTTCACTAGCTGAGCAACAA (SEQ ID NO: 9)
 B.anth 121 (SEQ ID NO: 10)

The *Bacillus anthracis* and *B. cereus* sasp-B sequence alignment showed a significant difference, namely a TAGCATT insert

BcerPub 1 AACAAAGCAACTTCTGGTGCTAGCATTCAAAGTACAAATGCTAGTTATGGTACAGAGTTT
 Banth 1G.....C.....
 BcerPub 61 TCAACTGAAACAGATGTACAAGCTGTAAAACAAGCAAACGCACAATCAGAAGCAAAGAAA
 Banth 61 G.G.....A.....A.....T.....
 BcerPub 121 GCACAAGCTTCTGGTGCA-----CAAAGTGCAAACGCTAGTTATGGTACAGAATTTGCA
 Banth 121 ..G.....TAGCATT.....CA....T.....
 BcerPub 175 ACTGAAACAGACGTGCATTCTGTGAAAAAACAAAATGCTAAGTCAGCTGCAAAACAA

(SEQ ID NO: 11)

Banth 181G.....AC.A.....

(SEQ ID NO: 12)

5 [37] Conclusion from above results are as follows: Only the *sasp-B* sequence from *Bacillus anthracis* diverges from that of its near neighbor, *B. cereus*, to any useful extent. The *sasp-2* sequences are identical, and the *sasp-1* sequences differ too little to be of use in distinguishing the two organisms.

[38] The underlined sequence TAGCATT (SEQ ID NO 107) represents an
10 insertion region useful for distinguishing *Bacillus anthracis* from other *Bacillus* species.

Example 4: Determination of optimal reaction conditions for maximal sensitivity

15 [39] By repeating the reaction condition optimization procedure described in Example 2 above, but upon a dilution series of *Bacillus anthracis* DNA down to 10 picograms per reaction, conditions were identified which resulted in maximum reaction sensitivity and specificity (as judged by the appearance of reaction products in ethidium bromide stained gel electrophoresis). By the criteria described, the best combination of conditions for use of BcSasp-B primers were as follows:

20 [40] In a 100 microliter reaction volume the final concentration of reactants were 10 millimolar tris buffer pH 8.3; 25 millimolar potassium chloride; 2 millimolar magnesium chloride; 0.2 millimolar each dinucleotide triphosphate dATP, dCTP, dGTP, and dTTP; 50 picomoles of each primer, 5 units of Taq polymerase (Perkin Elmer, Norwalk, Conn.). Thermalcycling conditions were the same as those described in Example 2 above.

25 [41] By viewing products with ethidium bromide stained gel electrophoresis it was determined that amplification specificity using these primers was quite good (only the expected product band and primers were visible), and product was visible down to 100 picograms per reaction.

30 Example 5: Addressing the necessary question: How conserved is the *sasp-B* sequence in *Bacillus anthracis*?

[42] In order to establish how conserved the promising stretch of *Bacillus anthracis* *sasp-B* gene sequence is, DNA from a variety of *B. anthracis* isolates was amplified and the amplicons sequenced. We acquired DNA from 38 geographically diverse anthrax isolates which was prepared by staff of the Centre for Applied Microbiological Research, Porton

Down, UK from the collection of Dr. Peter Turnbull. DNA from the 38 isolates was amplified using the *B. cereus* sasp-B primers and the (optimized) reaction conditions described in Example 4. The resulting PCR product was sequenced and analyzed as described above. The results presented in Figure 1 confirmed that not only is the six base insertion present in diverse isolates of *Bacillus anthracis*, but the sequence as a whole is quite well conserved.

[43] Referring now to Fig. 1A-C, there is illustrated the results of the CLUSTAL W alignment of 38 different *B. anthracis* strains, with the sequence of interest underlined in line 38. It is conserved in all strains. The table below sets forth the identification of the various strains used:

TABLE 1.

Legend				
<u>Bacteria</u>	<u>ASC#</u>	<u>NMRI#</u>	<u>Designation</u>	<u>Description or ATCC#</u>
<i>Bacillus anthracis</i>			Bapast	Institute Pasteur strain
<i>Bacillus anthracis</i>			Barec1	UM23, Thorne strain
<i>Bacillus anthracis</i>	152	1	NMRI#1	Namibia, 88
<i>Bacillus anthracis</i>	BA40D 2		NMRI#2	
<i>Bacillus anthracis</i>	92	4	NMRI#4	Zambia, hippo, and passaged mouse, 88
<i>Bacillus anthracis</i>	30	5	NMRI#5	Shropshire, acquired, 79
<i>Bacillus anthracis</i>	273	6	NMRI#6	XingJiang Province China, 92
<i>Bacillus anthracis</i>	63	10	NMRI#10	Etosha, soil via guinea pig, 86
<i>Bacillus anthracis</i>	BA42D 11		NMRI#11	
<i>Bacillus anthracis</i>	237	18	NMRI#18	Landkey, soil, mouse passage, 92
<i>Bacillus anthracis</i>	56	19	NMRI#19	Zimbabwe, human, 80
<i>Bacillus anthracis</i>	93	20	NMRI#20	Zambia, hippo, and passage guinea pig, 88
<i>Bacillus anthracis</i>		22	NMRI#22	MS191
<i>Bacillus anthracis</i>	91	23	NMRI#23	Zambia, hippo, 88
<i>Bacillus anthracis</i>	11	24	NMRI#24	NCTC5444, London, 28
<i>Bacillus anthracis</i>	64	25	NMRI#25	Russian vaccine, Sterne
<i>Bacillus anthracis</i>	29	26	NMRI#26	Waybridge, cow, traced to Senegal

<i>Bacillus anthracis</i>	238	28	NMRI#28	Landkey, soil, 92
<i>Bacillus anthracis</i>	245	32	NMRI#32	Sterne
<i>Bacillus anthracis</i>		35	NMRI#35	1+2+DMD
<i>Bacillus anthracis</i>	234	36	NMRI#36	Wessex, soil, 92
<i>Bacillus anthracis</i>		38	NMRI#38	Rvacc
<i>Bacillus anthracis</i>		39	NMRI#39	PR13P
<i>Bacillus anthracis</i>	264	40	NMRI#40	Zambia, contaminated soil, 91
<i>Bacillus anthracis</i>	43	41	NMRI#41	M36, passaged rabbits and rats
<i>Bacillus anthracis</i>	69	42	NMRI#42	New Hampshire, 57
<i>Bacillus anthracis</i>	65	43	NMRI#43	Brazil, cow, acquired, 82
<i>Bacillus anthracis</i>	192	50	NMRI#50	Landkey, soil
<i>Bacillus anthracis</i>	4	52	NMRI#52	M36, passaged in rabbits
<i>Bacillus anthracis</i>	3	53	NMRI#53	M36, derived from original challenge stock
<i>Bacillus anthracis</i>	2	54	NMRI#54	Griunard, 1950s
<i>Bacillus anthracis</i>	54	55	NMRI#55	Zimbabwe, human, 80
<i>Bacillus anthracis</i>		56	NMRI#56	1+2-
<i>Bacillus anthracis</i>	28	59	NMRI#59	Waybridge, cow, traced to Senegal, 78

NMRI is the Naval Medical Research Institute, Bethesda, Maryland

ATCC is the American Type Culture Collection

ASC is The Association Of Systematics Collections

5

Example 6: Scrutinizing the sequence of BcSasp-B primed amplicons from *Bacillus anthracis* near neighbors

[44] Following reports that other labs had mistakenly identified *B. thuringiensis* as *Bacillus anthracis*, 24 serotypes of *B. thuringiensis* were obtained in order to check whether

10 the sequence of sasp-B amplicons resembled those of other amplicons. DNA was prepared from *B. thuringiensis* as well as *B. cereus* liquid cultures following the method described by Zhou (Zhou, J., Bruns, M., and Jiedje, J. 1996. "DNA recovery from soils of diverse composition." Appl. Environ. Microbiol. 62:316-322), and amplified 100 nanograms of the DNA using *B. cereus* sasp-B primers. Product fragments were gel purified, extracted,

15 sequenced, and analyzed as described in Example 3 above. The resulting alignment

(Figure 2) includes *B. thuringiensis* and *B. cereus* sasp-B sequences as well as *Bacillus anthracis* sasp-B sequence from the previous example:

[45] Referring now to Figs. 2A, 2B, and 2C, the single *Bacillus anthracis* sequence (#37 which is the bottom row of Figs. 2A, 2B, & 2C) shows a unique pattern of sequence

5 divergence from the sasp-B sequence of these near neighbor isolates. The identities of the sequences are shown in Table 2 below:

10067613-020402

TABLE 2.

Legend:

<u>Bacteria</u>	<u>BGSC#</u>	<u>Serotype</u>	<u>Designation</u>
<i>Bacillus licheniformis</i> ,		5A2	5A2
<i>Bacillus thuringiensis</i>		4A1 serot-1	4A1
<i>Bacillus thuringiensis</i>		4A3 cry (thur) serot-1	4A3
<i>Bacillus thuringiensis</i>		4J2 aizawai, pacificus/serot-7	4J2
<i>Bacillus thuringiensis</i>	HD3	4B2 2 standard	BtB
<i>Bacillus thuringiensis</i>	HD4	4C3 3a standard	BtC
<i>Bacillus thuringiensis</i>	HD7	4E2 4a4b dendrolimus standard	BtE2
<i>Bacillus thuringiensis</i>		4E4 4a4b	BtE4
<i>Bacillus thuringiensis</i>	HD29	4G5 5a5b	BtG
<i>Bacillus thuringiensis</i>	HD10	4I1 6	BtI
<i>Bacillus thuringiensis</i>	HD11	4J4 7	BtJ
<i>Bacillus thuringiensis</i>	HD12	4K1 8 standard	BtK
<i>Bacillus thuringiensis</i>	HD537 4L3	9 standard	BtL
<i>Bacillus thuringiensis</i>	HD146 4M1	10 standard	BtM
<i>Bacillus thuringiensis</i>	HD201 4N1	11 antisera standard	BtM
<i>Bacillus thuringiensis</i>	HD542 4O1	12 standard	BtO
<i>Bacillus thuringiensis</i>	HD395	4P1 13 standard	BtP
<i>Bacillus thuringiensis</i>	ONR60A	4Q1 14	BtQ
<i>Bacillus thuringiensis</i>	HD511 4R1	15	BtR
<i>Bacillus thuringiensis</i>	HD521 4S2	16 standard	BtS
<i>Bacillus thuringiensis</i>	HD525 4T1	no flagellar antigen	BtT
<i>Bacillus thuringiensis</i>	HD541 4U1	11a11c	BtU
<i>Bacillus thuringiensis</i>		4V1 17	BtV
<i>Bacillus thuringiensis</i>	HD 867	4W1 18	BtW
<i>Bacillus thuringiensis</i>	IS720	4X1 21	BtX
<i>Bacillus thuringiensis</i>	HD868	4Y1 19 standard	BtY
<i>Bacillus thuringiensis</i>	HD501	4Z1 8a8c standard	BtZ
<i>Bacillus anthracis</i>	BA42D	11	NMRI#11
Unidentified <i>Bacillus</i>			003
Unidentified <i>Bacillus</i>		Taken from filled bag in "final mixing trailer"	1B
Unidentified <i>Bacillus</i>		Isolated from 1B culture as morphologically distinct colonies	1B/A
Unidentified <i>Bacillus</i>		Isolated from 25kg media drum, bentonite	III

		mixture	
Unidentified <i>Bacillus</i>		Isolated from bentonite spore stock	IV
<i>Bacillus cereus</i>	Genbank #M16813	NCBI Genbank database	Bcerpub
<i>Bacillus cereus</i>	ATCC 14579	Purchased from ATCC	Bcer1
<i>Bacillus cereus</i>	ATCC 11778	Purchased from ATCC	Bcer2
<i>Bacillus cereus</i>	ATCC 6464	Purchased from ATCC	Bcer3

BGSC is the *Bacillus* Genetic Stock Center, at The Ohio State University

[46] Based on the DNA sequence information in Figures 1 and 2, amino acid sequences were extrapolated and evaluated for the *sasp-B* genes from *Bacillus anthracis*, *Bacillus cereus* and *Bacillus thuringiensis*. These extrapolated sequences are shown below in an extrapolated amino acid sequence alignments for the *sasp-B* gene from *Bacillus anthracis*, *Bacillus cereus* and *Bacillus thuringiensis*. The identities of the sequences are shown in Table 3.

		15 16	30 31	45 46	60
1	4D4	NKATSGASIQSTNAS	YGTEFSTETDVQAVK	QANAQSEAKKAQASG	A-QSANASYGTEFA
2	Bcep	NKATSGASIQSTNAS	YGTEFSTETDVQAVK	QANAQSEAKKAQASG	A--QSANASYGTEFA
3	BtK	NKATSGASIQSTNAS	YGTEFATETNVQAVK	QANAQSEAKKAQASG	A--QSANASYGTEFA
4	BtB	NKATSGASIQSTNAS	YGTEFSTETDVQAVK	QANAQSEAKKAQASG	A--QSANASYGTEFA
5	Banth	NKATSGASIQSTNAS	YGTEFATETNVQAVK	QANAQSEAKKAQASG	ASIQSTNASYGTEFA
6	Bmyc	NKATSGASIQSTNAS	YGTEFATETNVQAVK	QANAQSEAKKAQASA	A--QSANASYGTEFA
		61	75 76		
1	4D4	TETDVHSVKKQNAKS	AAKQ (SEQ ID NO: 88)		
2	Bcep	TETDVHSVKKQNAKS	AAKQ (SEQ ID NO: 89)		
3	BtK	TETDVHAVKKQNAKS	AAKQ (SEQ ID NO: 90)		
4	BtB	TETDVHAVKKQNAQS	AAKQ (SEQ ID NO: 91)		
5	Banth	TETDVHAVKKQNAQS	AAKQ (SEQ ID NO: 92)		
6	Bmyc	TETDVHAVKKQNAQS	AAK (SEQ ID NO: 93)		

TABLE 3.

Legend:

Bacteria	BGSC#	Serotype	Designation
<i>Bacillus thuringiensis</i>	type strain		4D4
<i>Bacillus thuringiensis</i>	HD12	4K1 8 standard	BtK
<i>Bacillus thuringiensis</i>	HD3	4B2 2 standard	BtB
<i>Bacillus cereus</i>	published sequence	GenBank #M16813	Bcerp

5 Example 7: Design and evaluation of primers and probes to *Bacillus anthracis* specific sasp-B DNA sequence

[47] In the previous examples, the BcSasp-B primers were useful for evaluating the prevalence of the unique *Bacillus anthracis* sasp-B signature, but sequencing was required to distinguish amplicons of the several *Bacillus* species which could be amplified using the

10 *Bacillus cereus* primers. By studying the alignment of *Bacillus anthracis* and *Bacillus cereus* sasp-B sequences (above) potential anthracis specific primer and probe sites were identified (shown below, SEQ ID NO: 94 and 95). Eight oligonucleotides were designed with the aid of Oligo 4.0 and BLAST database search utilities then synthesized (all as described in Example 1 above) and evaluated experimentally in various combinations for their ability to
15 prime amplification of *Bacillus anthracis* only, using a panel of near neighbor *Bacillus* species. Three of the primer pairs were designed to incorporate the *Bacillus anthracis* insertion region into the three prime end of one primer per pair. This strategy greatly limited amplicon size and did not leave any *Bacillus anthracis* specific sequence for probe design.

[48] The combination of primers originally designated BaSPB7 and BaSPB8
20 (below) were sufficiently specific. From 100 nanograms *B. cereus* target a very faint product band of nearly (but not exactly) the correct size, was evident; when compared to signals from an amplified dilution series of *Bacillus anthracis* DNA, the signal from *Bacillus cereus* was approximately equivalent to product from 10 picograms - indicating 10,000 fold less efficient amplification. Bands were not visible at or near the correct size from products of *Bacillus*
25 *coagulans*, *Bacillus circulans*, *Bacillus globigii*, *Bacillus mycoides*, *Bacillus subtilis* or *Bacillus thuringiensis* amplification.

[49] In addition, these primers were for sequences flanking, rather than incorporating the *Bacillus anthracis* insertion region, thus leaving this region within the product for binding to probes designed to hybridize to this unique signature.

30

The *Bacillus anthracis* primer data (from analysis by Oligoprimer design software, National Biosciences, Plymouth, MN.) is summarized as follows:

BaSPB7 primer sequence:

5' GTT ATG GTA CAG AGT TTG CG 3' (SEQ ID NO: 94)

T_m = 57.4 °C (salt 1000.0 mM; oligo 0.6 pM)

T_d = 57.6 °C, G(25°C) = -34.7 kcal/mol, Mr = 6283

Ext. coeff.: 5.05 nmol/A260, 31.7 µg/A260

BaSPB8 primer sequence:

5' TTG TTT TGC AGC TGA TTG T 3' (SEQ ID NO: 95)

T_m = 58.3 °C (salt 1000.0 mM; oligo 0.6 pM)

T_d = 58.9 °C, G(25°C) = -34.1 kcal/mol, Mr = 5911

Ext. coeff.: 5.82 nmol/A260, 34.4 µg/A260

[50] Optimal amplification conditions were determined in the same manner described in Examples 2 and 4 above. Optimal amplification conditions were identified as follows:

[51] Thermalcycling: Amplifications were performed in a Perkin Elmer 9600 thermocycler with the following thermal cycling regime: 94oC for 5 minutes, then 40 repeating cycles of 94oC for 30 seconds, 50oC for 30 seconds and 72oC for 30 seconds, followed by a 7 minute 72oC final extension step.

[52] Reaction mixture: Each 100ul reaction contained 0.1 millimolar each dATP, dCTP, dGTP and dTTP, 25 picomoles each primer, 10 millimolar Tris-HCl pH 8.3, 2 millimolar MgCl₂, 25 millimolar KCl, 2.5 units of Taq polymerase (Perkin Elmers, Norwalk, Conn.) and 100 ng or less of template DNA.

The uniqueness of these primers may be seen by a *Bacillus anthracis* and *Bacillus cereus* sasp-B sequence alignment emphasizing *Bacillus anthracis* specific primer sequences:

BcerPub	1	AACAAAGCAACTTCTGGTGCTAGCATTCAAAGTACAAATGCTAGTTATGGTACAGAGTTT
Banth	1G.....C..... <u>GTTATGGTACAGAGTTT</u>
		--primer BaSPB7--

BcerPub 61 TCAACTGAAACAGATGTACAAGCTGTAAAACAAGCAAACGCACAATCAGAAGCAAAGAAA
 Banth 61 GCG.....A.....A.....T.....
 -->

BcerPub 121 GCACAAGCTTCTGGTGCA-----CAAAGTGCAAACGCTAGTTATGGTACAGAATTTGCA
 Banth 121 ..G.....TAGCATT.....CA....T.....

BcerPub 175 ACTGAAACAGACGTGCATTCTGTGAAAAACAAAATGCTAAGTCAGCTGCAAAACAAA

(SEQ ID NO: 96)

Banth 181G.....ACAATCAGCTGCAAAACAAA

(SEQ ID NO: 97)

<--primer BaSPB8-

[53] The *Bacillus anthracis* and *Bacillus cereus* sasp-B sequence alignment shows the sequence similarity between the present *Bacillus anthracis* sasp B DNA (which is a 240 base pair amplicon as described above) and the corresponding region of its most similar known sequence, the *Bacillus cereus* sasp B gene. This alignment, run in the CLUSTALW program described herein, yields a similarity score of 89%, using default parameters. As is known in the art, the default parameters for nucleic acid pairwise alignments are gap opening penalty = 15; gap extension penalty = 6.66. The IUB matching protocol is used -All matches score 1.9; all mismatches for IUB symbols score 0. The CLUSTAL matching protocol can also be used. In this case, matches score 1.0 and mismatches score 0.0. In either case, CLUSTALW comparison of the *Bacillus anthracis* sasp B DNA and the comparable *Bacillus cereus* sasp B DNA yields a score of 89%.

[54] Similarly, given the present *Bacillus anthracis* sasp B DNA, a similarity score from the NCBI GenBank nucleotide database using BLAST (default parameters, version 2.2.1) is easily obtained. As is known in the art, the BLAST defaults for nucleotide sequences are: -3 for a nucleotide mismatch; +1 for a nucleotide match; and 0 gap penalty.

The matrix used by default is Blosum62.

[55] In the present case, a BLAST search revealed that the closest sequence corresponds to the *Bacillus cereus* small acid-soluble spore protein, Accession Number M16813. This corresponds with the discussion in this Example. Using BLAST parameters, the two nucleotide sequences have an identity of 86%.

[56] A manual analysis of the *Bacillus anthracis* and *Bacillus cereus* sasp-B-sequence alignment yields a similarity score of 90-91%. To perform a manual analysis, one

of skill in the art would add the number of nucleotides that differ among the two sequences. One of skill would then subtract that number from the total number of nucleotides in the sasp-B *Bacillus anthracis* sequence and divide the resulting number by the total number of nucleotides in the sasp-B *Bacillus anthracis* sequence. The resulting number multiplied by 100 yields the similarity score.

[57] Accordingly, those of skill in the art would recognize that the Sasp-B DNA of *Bacillus anthracis* is homologous but not identical to that of *Bacillus cereus*. Thus, this invention includes any sasp-B from *Bacillus anthracis* that might include minor single base differences (polymorphisms) from SEQ ID NO: 97 (or identical SEQ ID NO:87), yet, maintain the insert of SEQ ID NO: 107.

Example 8: Selection and evaluation of probes for detection of *Bacillus anthracis* sasp-B amplicons

[58] Three of the oligonucleotides evaluated as primers incorporated the *Bacillus anthracis* specific insertion region, and having designed primers flanking the insertion region, these oligos were tested as probes to confirm the identity of the amplicons; only amplicons from *B. anthracis* would include the 6 base insertion, as follows:

Alignment of *Bacillus cereus* and *Bacillus anthracis* Sasp-B sequences emphasizing probed locations

Bcer	AACAAAGCAACTTCTGGTGCTAGCATTCAAAGTACAAATGC
Banth	AACAAGGCAACTTCTGGTGCTAGCATTCAAAGCACAAATGC
Bcer	TAGTTATGGTACAGAGTTTTCAACTGAAACAGATGTACAAGCTGTAAAACAAGCAAACGCACAA
Banth	TAGTTATGGTACAGAGTTTGC GACTGAAACAAATGTACAAGCAGTAAAAAAGCAAACGCACAAT
Bcer	TCAGAAGCAAAGAAAGCACAAAGCTTCTGGTGCA-----AAAGTGCAAACGCTAGTTATGGTACAGAATTTGCAA
Banth	CAGAAGCTAAGAAAGCGCAAGCTTCTGGTGCTAGCATTCAAAGCACAAATGCTTTGCATAGTTATGGTACAGAAA

	↑ location of probes tested ↑
Bcer	CTGAAACAGACGTGCATTCTGTGAAAAAACAAATGCTAAGTCAGCTGCAAAACAA (SEQ ID NO: 98)
Banth	CTGAAACAGACGTGCATGCTGTGAAAAAACAAATGCACAATCAGCTGCAAAACAA (SEQ ID NO: 99)

Three oligonucleotides evaluated for use as *Bacillus anthracis* probes:

BaSPB2: (inverted, 'lower strand' sequence): 5' GCATTTGTGCTTTGAATGCTA 3' (SEQ ID NO: 100)

5 BaSPB4: (inverted, 'lower strand' sequence): 5' CATTGTGCTTTGAATGCTA 3' (SEQ ID NO: 101)

BaSPB5: (direct, 'upper strand' sequence): 5' AGCTTCTGGTGCTAGCATT 3' (SEQ ID NO: 102)

[59] Oligos (shown above) were tested as probes in the following manner:

[60] *Bacillus anthracis* DNA (100 nanograms per 100 microliter reaction of the
10 Sterne strain) was amplified using Biotin labeled BaSPB7 and BaSPB8 primers which had
been synthesized to our order by Life Technologies (Gaithersburg, MD).

[61] Following the instructions for the Universal GeneComb Kit (Bio-Rad,
Richmond, CA), 8 picomoles and 5 picomole quantities of each oligo (henceforth called
probe) in the kit binding buffer was spotted on the GeneComb test kit card nitrocellulose
15 surface, and bound to the nitrocellulose using the UV Stratalinker 1800 (Stratagene, La Jolla,
CA) set to auto crosslink mode. The biotinylated amplicon was denatured, and allowed to
migrate across the (bound) probe spots, hybridizing to them in the process. The resulting
bound biotinylated hybrid was then reacted with the kit streptavidin/alkaline phosphatase
conjugate followed by kit chromogenic substrate-enabling visualization of the
20 probe/amplicon hybrids. Evaluation of the test spot signal intensity guided the choice of
probe for further evaluation. BaSPB4 proved most sensitive in this test, so succeeding work
was done with this probe.

[62] In order to roughly gauge sensitivity of the system, amplification was
performed using Biotinylated BaSPB7 and BaSPB8 primers, the target(s) being a dilution
25 series of *Bacillus anthracis* DNA (Sterne strain) whose concentration (prior to dilution) had
been carefully determined using the Beckman DU 640 Spectrophotometer. The dilutions
amplified were (total input per 100 microliter reaction): 10, 5, and 1 picogram(s), 100, 50,
and 20 femtograms. Following the GeneComb test kit instructions, one tenth of each PCR
was reacted with (bound) BaSPB4 and the resulting hybrids 'developed'. The developed test
30 strips reflected the input DNA dilutions in color intensity, with even the 20 femtogram
reaction yielding a visible spot. (Data not shown.) The results comprised a visible spot for
each dilution tested (10 pg down to 20 fg), with the negative PCR and kit control showing no
spots.

[63] Finally, probe BaSPB4 was tested for specificity. Fifty nanograms of DNA
35 from each of the following *Bacillus* species was amplified using biotinylated BaSPB7 and

BaSPB8: *Bacillus anthracis*, *Bacillus thuringiensis*, *Bacillus cereus*, *Bacillus mycoides*, *B. subtilis*, *Bacillus globigii*. The denatured amplicon of each DNA species was reacted against the bound BaSPB4 probe and the test strips developed. Only the *Bacillus anthracis* amplicon resulted in any signal (which was quite intense); none of the other species bound to the probe in order to result in a signal. Conclusion: BaSPB4 binds *Bacillus anthracis* DNA specifically. BaSPB4 binding specificity was demonstrated with the BioRad universal GeneComb System (data not shown). 100 ng of each species of DNA amplified was placed on each panel, as follows (1 - 8):

Species Amplified:

- 1) *Bacillus anthracis*
- 2) *Bacillus thuringiensis*
- 3) *Bacillus cereus*
- 4) *Bacillus mycoides*
- 5) *Bacillus subtilis*
- 6) *Bacillus globigii*
- 7) Negative PCR
- 8) Kit positive control

[64] The *Bacillus anthracis* showed a large spot; the other panels were blank, except for the positive control. This showed that only the amplified *Bacillus anthracis* sasp B sequence reacted with the probe.

[65] Some potential detection systems for the *Bacillus anthracis* specific primer/probe system are as follows:

[66] The primer/probe system described above is ideally suited to the 5'nuclease fluorescence homogeneous assay in which the accumulation of specific amplicon is monitored as fluorescence is released from the probe by Taq polymerase during the amplification of target DNA to which the probe anneals. This system is described in U.S. patents 5,538,848; 5,723,591; and 5, 876,930, hereby incorporated by reference.

[67] The primer/probe system described is also suited to amplification followed by separate hybridization of biotinylated amplicon molecules to bound probe in a colormetric microwell plate type assay (J. Mulder, N. McKinney, C. Christopherson, J. Sninsky, L. Greenfield and S. Kwok. Rapid and Simple PCR Assay for Quantitation of HIV-I RNA in Plasma: Applications to Acute Retroviral Infection. J. Clin. Micro. 37(2): 292-300, 1994);

or, in similar manner, as with the automated AmpliCore integrated PCR+detection devices (Roche, Pleasanton, CA), and as described, a simple dot blot type assay.

[68] There is an increasing number of PCR devices and coordinated detection strategies; the primers/probes and amplification system described are robust, specific, and sensitive enough to be adapted to most of these.

[69] Regardless of detection format, the described assay could be used to monitor the presence of anthrax in the environment (such as for investigation by military and law enforcement agencies of clandestine production of anthrax for illicit use; for Public Health and law enforcement agencies to test suspicious spore-like powders; to check for anthrax in suspected cases of bio-terrorist attack).

[70] This assay is also well suited as a rapid, specific, and sensitive method for detecting anthrax in biological fluids such as blood, sputum, and feces in clinical and Public Health labs, as well in the field, and for autopsies.

Example 9: Development of *Bacillus globigii* specific PCR based on *sasp* gene sequence

[71] In the same manner as described in Example 1 above, a *sasp* gene (*sasp*-gamma in this case) sequence was identified for the production of primers specific for *Bacillus globigii* sequence. Primers and amplification conditions were designed (see Figure 3) for heterologous PCR based on published sequence for the *Bacillus subtilis* *sasp* E gene (*sasp*-gamma) acquired from GenBank (accession number M16184). After sequencing amplicons from *Bacillus globigii* (generated using the *Bacillus subtilis* primers), and aligning *Bacillus globigii* sequence with the published *Bacillus subtilis* sequence, *Bacillus globigii* specific primers were designed taking advantage of the differences in the sequence. After searching the databases to be sure that the new *Bacillus globigii* primers were not homologous to other sequences, and optimizing amplification conditions, a panel of *Bacillus* species were amplified to check primer specificity. Amplicons of the correct size were produced only from *Bacillus* designated as *Bacillus globigii*, for all but the most arcane intents and purposes (there is disagreement among a very few researcher as to whether *Bacillus subtilis niger* and *Bacillus atrophaeus* are, in fact, genetically different from *Bacillus globigii* at all); importantly, the new primers did not amplify *Bacillus subtilis* or *Bacillus amyloliquifaciens* -which are distinct species, yet very closely related to *Bacillus globigii*. Referring now to Fig. 3, there is shown an alignment of *Bacillus subtilis* *sasp*-gamma sequence (from Genbank) (Bs_pub_SSPE) with *Bacillus globigii* sequence (upper strand)

showing the location of the primer sequences and how their sequence compares to the known *Bacillus subtilis* sequence.

[72] The BgSaspGam primers produce *B. globigii* specific PCR product, as was demonstrated in an Nuseive-Agarose gel (data not shown). The gel showed approximately a 135b *Bacillus globigii* specific amplicon. No amplification of negative controls in *Bacillus cereus*; *Bacillus amyoliquifaciens*, *Bacillus megaterium*, or *Bacillus globisporus* was observed. Amplification was observed with *Bacillus atrophaeus* (ATCC 6455 and 49337) and *Bacillus niger*. It should be noted that *Bacillus subtilis niger* and *Bacillus atrophaeus* have been officially designated *Bacillus globigii* since they are virtually indistinguishable from *Bacillus globigii* at the molecular level. Near neighbors *Bacillus subtilis*, *Bacillus globisporus* and *Bacillus megatarium* do not amplify with the BgSaspGam primers.

Bacillus globigii sasp-gamma primers:

BgSaspGam 5' 5' ACATGGCTAACTCAAACAACAA 3' (SEQ ID NO: 103)

BgSaspGam 3' 5' GGTTTGTCTTCTTACTTGTTGTAC 3' (SEQ ID NO: 104)

[73] Reaction conditions successfully employed using the above primers are as follows:

[74] Reaction mixture composition: In a 100 microliter reaction volume the final concentration of reactants were 10 millimolar tris buffer pH 8.3; 25 millimolar potassium chloride; 2 millimolar magnesium chloride; 0.2 millimolar each dinucleotide triphosphate dATP, dCTP, dGTP, and dTTP; 50 picomoles of each primer, 5 units of Taq polymerase (Perkin Elmer, Norwalk, Conn.).

[75] Thermalcycling conditions using TC9600 (Perkin Elmer, Norwalk, Conn.): A 5 minute 94 degree C. initial denaturation step was followed by 40 three step cycles of 94 degrees C. for 30 sec., 50 degrees C. for 30 sec., 72 degrees C. for 30 sec. A final extension step of 7 minutes at 72 degrees C. completed the thermalcycling.

[76] There a number of potential uses of the research *Bacillus globigii* specific amplification system. *Bacillus globigii* is used as a nonpathogenic surrogate, replacing *Bacillus anthracis*, for purposes of modeling aerosol spore distribution under various environmental conditions, as well as for testing spore collection hardware. The agencies carrying out these endeavors have had trouble finding a way of detecting only the *Bacillus globigii* used in their experiments; detection systems have been non-specific, resulting in false alarms and compromised data.

Example 10: Development of *Clostridium perfringens* PCR based on sasp gene sequence

[77] In a manner similar to the above descriptions, a sasp gene (sasp-2 in this case) sequence was identified for the production of primers for amplification of *Clostridium perfringens* sequence. Primers and amplification conditions were designed and carried out using *Clostridium perfringens* DNA. While amplification successfully produced product of the correct size (when viewed by ethidium bromide gel electrophoresis), near neighbor DNA has yet to be evaluated in order to assess specificity of these primers.

Clostridium perfringens sasp-2 primers:

CPssp2-1: 5' AATAACTAAGGAGGAATGAAAAATGT 3' (SEQ ID NO:105)

Cpssp2-2: 5' TTGTTCTACCATTCTTTTAACCATT 3' (SEQ ID NO: 106)

[78] The following reaction conditions were successfully employed using the above primers:

[79] Reaction mixture composition: In a 100 microliter reaction volume the final concentration of reactants were 10 millimolar tris buffer pH 8.3; 25 millimolar potassium chloride; 2 millimolar magnesium chloride; 0.2 millimolar EACH dinucleotide triphosphate dATP, dCTP, dGTP, and dTTP; 50 picomoles of each primer, 5 units of Taq polymerase (Perkin Elmer, Norwalk, Conn.).

[80] Thermalcycling conditions using TC9600 (Perkin Elmer, Norwalk, Conn.): A 5 minute 94 degree C. initial denaturation step was followed by 40 three step cycles of 94 degrees C. for 30 sec., 50 degrees C. for 30 sec., 72 degrees C. for 30 sec. A final extension step of 7 minutes at 72 degrees C. completed the thermalcycling.

[81] There are a number of potential uses of *Clostridium perfringens* specific PCR. *Clostridium perfringens* is officially listed as a biological weapon agent, so uses would be similar to those described for the *Bacillus anthracis* specific primers.

[82] Also, using these primers for heterologous PCR of *Clostridium botulinum* (a serious health threat and potential biological weapon) in order to acquire sequence information for the design of primers specific to that organism. Such primers, and any probe so identified in the process, would also be useful in the same manner described for the *Bacillus anthracis* specific detection research system described above.

[83] Having described the present invention, it will be apparent that other embodiments are possible in light of the present teachings. For example, other DNA amplification methods

besides PCR are known, such as the Q-beta replicase method. Certain of these methods may be used in a single-step amplification/detection protocol, based, for example, on the unique *Bacillus anthracis* sasp-B insertion TAGCATT (SEQ ID NO 107).

[84] Accordingly, the present invention should be understood to encompass subject matter limited only by the scope of the appended claims. Furthermore, having described the amino acid structure of at least a portion of the *B. anthracis* sasp-B gene, tests which directly test the sasp-B protein are now possible.

Example 11: The production of antibodies to sasp polypeptides.

[85] The novel sasp polypeptides of the invention can also be used to produce antibodies which are specifically immunoreactive or bind to epitopes of the sasp polypeptides. Antibodies of the invention specifically include antibodies which bind to unique polypeptides produced by the *B. anthracis* DNA sequence shown in Example 6 (and translated in Example #5) and identified as NMRI#11.

[86] The term "antibody" as used in this invention includes intact molecules as well as fragments thereof which are capable of binding the epitopic determinant. These antibody fragments retain some ability to selectively bind with its antigen or receptor and are defined as follows: Fab, the fragment which contains a monovalent antigen-binding fragment of an antibody molecule, can be produced by digestion of whole antibody with the enzyme papain to yield an intact light chain and a portion of one heavy chain; Fab', the fragment of an antibody molecule can be obtained by treating whole antibody with pepsin, followed by reduction, to yield an intact light chain and a portion of the heavy chain; two Fab' fragments are obtained per antibody molecule; F(ab').sub.2, the fragment of the antibody that can be obtained by treating whole antibody with the enzyme pepsin without subsequent reduction; F(ab').sub.2 is a dimer of two Fab' fragments held together by two disulfide bonds; Fv, defined as a genetically engineered fragment containing the variable region of the light chain and the variable region of the heavy chain expressed as two chains; and single chain antibody ("SCA"), defined as a genetically engineered molecule containing the variable region of the light chain, the variable region of the heavy chain, linked by a suitable polypeptide linker as a genetically fused single chain molecule.

[87] Methods of making these fragments are known in the art. (See for example, Harlow and Lane, Antibodies: A Laboratory Manual, Cold Spring Harbor Laboratory, New York (1988), incorporated herein by reference).

[88] As used in this invention, the term "epitope" means any antigenic determinant on an antigen to which the paratope of an antibody binds. Epitopic determinants usually consist of chemically active surface groupings of molecules such as amino acids or sugar side-chains and usually have specific three-dimensional structural characteristics, as well as specific charge characteristics. The unique amino acid sequence of the present sasp polypeptides provides a correspondingly unique epitope in three-dimensional space.

[89] Monoclonal antibodies are made from antigen containing fragments of the unique polypeptide by methods well known in the art (Kohler, et al., Nature, 256:495, 1975; Current Protocols in Molecular Biology, Ausubel, et al., ed., 1989).

[90] Monoclonal or polyclonal antibodies which bind to the sasp polypeptide of the invention can be prepared using an intact polypeptide or fragments containing small peptides of interest as the immunizing antigen. The polypeptide or a *B. anthracis* DNA sequence as shown in Example 6 and identified as NMRI#1, used to immunize an animal can be derived from translated cDNA or chemical synthesis which can be conjugated to a carrier protein, if desired. Such commonly used carriers, which are chemically coupled to the peptide, include keyhole limpet hemocyanin (KLH), thyroglobulin, bovine serum albumin (BSA), and tetanus toxoid. The coupled peptide is then used to immunize the animal (e.g., a mouse, a rat, or a rabbit).

[91] If desired, polyclonal or monoclonal antibodies can be further purified, for example, by binding to and elution from a matrix to which the polypeptide, or a peptide to which the antibodies were raised, is bound. Those of skill in the art will know of various techniques common in the immunology arts for purification and/or concentration of polyclonal antibodies, as well as monoclonal antibodies (See for example, Coligan, et al., Unit 9, Current Protocols in Immunology, Wiley Interscience, 1991, incorporated by reference).

[92] It is also possible to use the anti-idiotypic technology to produce monoclonal antibodies which mimic an epitope. For example, an anti-idiotypic monoclonal antibody made to a first monoclonal antibody will have a binding domain in the hypervariable region which is the "image" of the epitope bound by the first monoclonal antibody. Thus, in the present invention, an anti-idiotypic antibody produced from an antibody which binds to the synthetic peptide of the invention can bind to the site on *B. anthracis* sasp which forms the spore, thereby preventing sasp from participating in spore assembly.

[93] Polynucleotide sequences encoding the polypeptide or synthetic peptide (*B. anthracis* DNA sequence shown in Example 6 and identified as NMRI#11) of the invention

can be expressed in either prokaryotes or eukaryotes. Hosts can include microbial, yeast, insect and mammalian organisms. Methods of expressing DNA sequences having eukaryotic or viral sequences in prokaryotes are well known in the art. Biologically functional viral and plasmid DNA vectors capable of expression and replication in a host are known in the art.

- 5 Such vectors are used to incorporate DNA sequences of the invention.

[94] Accordingly, the present invention should be understood to encompass subject matter limited only by the scope of the appended claims.

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